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(FILE 'HOME' ENTERED AT 18:28:32 ON 13 JUN 2004)

FILE 'MEDLINE, BIOSIS' ENTERED AT 18:28:48 ON 13 JUN 2004

L1           0 S PRO7170  
L2        3446 S GLUCOSE (L) FATTY ACID? (L) UPTAKE  
L3       17962 S (GLUCOSE OR FATTY ACID?) (W) UPTAKE  
L4       468 S GLUCOSE (L) FATTY ACID? (L) UPTAKE (L) SKELETAL MUSCLE  
L5      277 DUP REM L4 (191 DUPLICATES REMOVED)  
L6      238 S L5 AND PY<2003  
L7      386 S (ASHKENAZI, A?)/AU  
L8       0 S L6 AND L7  
L9       0 S EXMAD

XX PRO polynucleotides used to produce polypeptides used to target  
 PT bioactive molecules such as toxins, radiolabels or antibodies, to  
 PT specific cells, to cause targeted cell death.  
 XX  
 PS Claim 12: FIG 326; 935pp; English.

CC The present invention describes human secreted and transmembrane PRO  
 CC proteins. The PRO proteins have cytostatic activity. The PRO proteins  
 CC can be used for targeted delivery of bioactive molecules, such as  
 CC toxins, radiolabels or antibodies, that cause cell death. PRO nucleotide  
 CC sequences, and their fragments, can be used as hybridisation probes, in  
 CC chromosomal and gene mapping, and in the generation of anti-sense RNA  
 CC and DNA. They may also be used to produce transgenic animals which are  
 CC used to develop and screen therapeutically useful reagents. The PRO  
 CC nucleotide and protein sequence can be used for tissue typing and in  
 CC treating cancer. Anti-PRO antibodies can be used in diagnostic assays.  
 CC AAF4470 to AAF4470 represent PCR primers. AAF44087 to AAF44269 and  
 CC AAB65154 to AAB65300 represent human PRO polynucleotide and protein  
 CC sequences given in the exemplification of the present invention.  
 XX Sequence 482 AA;

Query Match 100.0%; Score 2429; DB 22; Length 482;

Best Local Similarity 100.0%; Pred. No. 1e-148;  
Matches 482; Conservative 0; Mismatches 0; Indels 0; Gaps 0;Qy 1 MGCLMGLALPFFCMEVGYSSAGPSPSTRADMTTDDTEVAMTLRPGHALETQL 60  
Db 1 MGCLMGLALPFFCMEVGYSSAGPSPSTRADMTTDDTEVAMTLRPGHALETQL 60  
Qy 61 SAETSSRASPTPAGPIPEAETRGAKRISPARETRSFTKTPSPNEMYLATSVETAAAGSPE 120  
Db 61 SAETSSRASPTPAGPIPEAETRGAKRISPARETRSFTKTPSPNEMYLATSVETAAAGSPE 120  
Qy 121 GAGMTTVQITGSDEEAFLFDTLCTDDSEBEEAKTLMDLTLAHTSTEAKGLSSESSAS 180  
Db 121 GAGMTTVQITGSDEEAFLFDTLCTDDSEBEEAKTLMDLTLAHTSTEAKGLSSESSAS 180  
Qy 181 DGPHPVITPERASESSASGDGPHEVITPERASESSASGDGPHEVITPWSGPGDVTLAE 240  
Db 181 DGPHPVITPERASESSASGDGPHEVITPERASESSASGDGPHEVITPWSGPSSDVTLLAE 240  
Qy 242 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
Db 241 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
Qy 301 TEVTAETLSTAGTTESSAAPHATGTPLPNTSATEREYTAGTTLSGALTVVSNPLE 360  
Db 301 TEVTAETLSTAGTTESSAAPHATGTPLPNTSATEREYTAGTTLSGALTVVSNPLE 360  
Qy 367 ETSALSVEPESYVKVSGAPVSIEAGSAVGKTTSFAGSSASSYSPSEALKNTPSETPT 420  
Db 361 ETSALSVEPESYVKVSGAPVSIEAGSAVGKTTSFAGSSASSYSPSEALKNTPSETPT 420  
Qy 421 MDIATKGPPPTSQRPLPSPPPTTNSRCRNTSLAKITTSAXTMKQQPRPLPGRGP 480  
Db 421 MDIATKGPPPTSQRPLPSPPPTTNSRCRNTSLAKITTSAXTMKQQPRPLPGRGP 480  
Qy 481 QT 482  
Db 481 QT 482

DE Human EXMAD-3 SEQ ID NO: 3.  
 XX KW Extracellular matrix and adhesion-associated protein; EXMAD; cancer;  
 KW inflammation; reproductive disorder; cardiovascular disorder;  
 KW immune disorder; musculoskeletal disorder; developmental disorder;  
 KW gastrointestinal disorder; cell proliferation disorder;  
 XX OS Homo sapiens.  
 XX WO200608380-A2.  
 XX PD 16-NOV-2000.  
 XX PF 10-MAY-2000; 20000WO-US12811.  
 XX PR 11-MAY-1999; 99US-013343.  
 XX PR 23-AUG-1999; 99US-0150409.  
 XX PA (INCYT) INCYTE GENOMICS INC.  
 XX PA Bandman O, Hillman JL, Tang YT, Lal P, Yue H, Baughn MR, Lu DAM;  
 PI Azimzai Y;  
 XX DR WPI; 2001-007395/01.  
 XX DR N-PSDB; AAC66892.  
 XX PT Isolated polynucleotide encoding extracellular matrix or  
 PT adhesion-associated protein (EXMAD) useful for diagnosing, treating, or  
 PT preventing disorders associated with expression of EXMAD such as  
 PT proliferative, immune and genetic disorders -  
 XX PS Claim 1; Page 89-90; 129pp; English.  
 XX  
 CC The present invention provides the protein and coding sequences for 25  
 CC novel extracellular matrix and adhesion-associated proteins (EXMADS).  
 CC These are designated EXMAD-1, EXMAD-2, EXMAD-3, EXMAD-4,  
 CC EXMAD-5, EXMAD-6, EXMAD-7, EXMAD-8, EXMAD-9, EXMAD-10, EXMAD-11, EXMAD-12,  
 CC EXMAD-13, EXMAD-14, EXMAD-15, EXMAD-16, EXMAD-17, EXMAD-18, EXMAD-19,  
 CC EXMAD-20, EXMAD-21, EXMAD-22, EXMAD-23, EXMAD-24 and EXMAD-25. They are  
 CC useful in the prevention and treatment of cancer, cell proliferation and -  
 CC cardiovascular, reproductive, immune, musculoskeletal, developmental and -  
 XX SQ Sequence 482 AA;  
 Query Match 100.0%; Score 2429; DB 22; Length 482;  
 Best Local Similarity 100.0%; Pred. No. 1e-148;  
 Matches 482; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Qy 1 MGCLMGLALPFFCMEVGYSSAGPSPSTRADMTTDDTEVAMTLRPGHALETQL 60  
 Db 1 MGCLMGLALPFFCMEVGYSSAGPSPSTRADMTTDDTEVAMTLRPGHALETQL 60  
 Qy 61 SAETSSRASPTPAGPIPEAETRGAKRISPARETRSFTKTPSPNEMYLATSVETAAAGSPE 120  
 Db 61 SAETSSRASPTPAGPIPEAETRGAKRISPARETRSFTKTPSPNEMYLATSVETAAAGSPE 120  
 Qy 121 GAGMTTVQITGSDEEAFLFDTLCTDDSEBEEAKTLMDLTLAHTSTEAKGLSSESSAS 180  
 Db 121 GAGMTTVQITGSDEEAFLFDTLCTDDSEBEEAKTLMDLTLAHTSTEAKGLSSESSAS 180  
 Qy 181 DGPHPVITPERASESSASGDGPHEVITPERASESSASGDGPHEVITPWSGPGDVTLAE 240  
 Db 181 DGPHPVITPERASESSASGDGPHEVITPERASESSASGDGPHEVITPWSGPSSDVTLLAE 240  
 Qy 241 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
 Db 241 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
 Qy 301 TEVTAETLSTAGTTESSAAPHATGTPLPNTSATEREYTAGTTLSGALTVVSNPLE 360  
 Db 301 TEVTAETLSTAGTTESSAAPHATGTPLPNTSATEREYTAGTTLSGALTVVSNPLE 360  
 Qy 367 ETSALSVEPESYVKVSGAPVSIEAGSAVGKTTSFAGSSASSYSPSEALKNTPSETPT 420  
 Db 361 ETSALSVEPESYVKVSGAPVSIEAGSAVGKTTSFAGSSASSYSPSEALKNTPSETPT 420  
 Qy 421 MDIATKGPPPTSQRPLPSPPPTTNSRCRNTSLAKITTSAXTMKQQPRPLPGRGP 480  
 Db 421 MDIATKGPPPTSQRPLPSPPPTTNSRCRNTSLAKITTSAXTMKQQPRPLPGRGP 480  
 Qy 481 QT 482  
 Db 481 QT 482

Qy 181 DGPHPVITPERASESSASGDGPHEVITPWSGPGDVTLAE 240  
 Db 181 DGPHPVITPERASESSASGDGPHEVITPWSGPSSDVTLLAE 240  
 Qy 241 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
 Db 241 ALVTVTNIVINCSITEETTSS1PAGSDIDLIPTEGVKASSTSDDPALPDSTEAKPHI 300  
 AC AAB27225;  
 XX 27-MAR-2001 (First entry)  
 XX DT Tue Jan 13 09:40:37 2004

RESULT 4  
 AAB27225  
 ID AAB27225 standard; Protein; 482 AA.  
 AC AAB27225;  
 XX 27-MAR-2001 (First entry)

Tue Jan

Qy 361 ETSALSVETPSVYKVSAAVPVSIEAGSAGVKTTSFAGSSASSYSPSEALNFTPSEPT 420  
 Db 361 ETSALSVETPSVYKVSAAVPVSIEAGSAGVKTTSFAGSSASSYSPSEALNFTPSEPT 420  
 PR 24-AUG-2000; 2000WO-US23328.  
 PR 01-DEC-2000; 2000WO-US32658.  
 PR 28-FEB-2001; 2001WO-US06500.  
 PR 30-MAY-2001; 2001WO-US17443.  
 PR 01-JUN-2001; 2001WO-US17800.  
 PR 20-JUN-2001; 2001WO-US19622.  
 PR 29-JUN-2001; 2001WO-US21066.  
 PR 09-JUL-2001; 2001WO-US21755.  
 PR 26-AUG-1997; 97US-056971P.  
 PR 17-SEP-1997; 97US-059115P.  
 PR 18-SEP-1997; 97US-059263P.  
 PR 19-SEP-1997; 97US-059583P.  
 PR 27-OCT-1997; 97US-063329P.  
 PR 24-OCT-1997; 97US-06285P.  
 PR 24-OCT-1997; 97US-062816P.  
 PR 27-OCT-1997; 97US-063082P.  
 PR 29-OCT-1997; 97US-063733P.  
 PR 21-NOV-1997; 97US-063640P.  
 PR 25-NOV-1997; 97US-066840P.  
 PR 16-DEC-1997; 97US-069694P.  
 PR 09-FEB-1998; 98US-074086P.  
 PR 09-FEB-1998; 98US-074092P.  
 PR 25-MAR-1998; 98US-079294P.  
 PR 08-APR-1998; 98US-081049P.  
 PR 10-AUG-1998; 98US-05998P.  
 PR 18-AUG-1998; 98US-07000P.  
 PR 09-SEP-1998; 98US-089601P.  
 PR 10-SEP-1998; 98US-089803P.  
 PR 09-FEB-1998; 98US-09811P.  
 PR 10-SEP-1998; 98US-09812P.  
 PR 17-SEP-1998; 98US-10858P.  
 PR 24-SEP-1998; 98US-101922P.  
 PR 28-OCT-1998; 98US-106032P.  
 PR 20-NOV-1998; 98US-109304P.  
 PR 23-MAR-1999; 99US-115778P.  
 PR 15-JUN-1999; 99US-13965P.  
 PR 20-JUL-1999; 99US-145070P.  
 PR 26-JUL-1999; 99US-14698P.  
 PR 17-AUG-1999; 99US-14936P.  
 PR 07-DEC-1999; 99US-169495P.  
 PR 15-NOV-2001; 2001US-0002799.  
 XX (GETH ) GENENTECH INC.

XX PD 28-NOV-2002.  
 XX PF 01-FEB-2002; 2002US-0066500.  
 XX PR 14-JUL-1998; 98WO-US14552.  
 PR 10-SEP-1998; 98WO-US18824.  
 PR 14-SEP-1998; 98WO-US19093.  
 PR 16-SEP-1998; 98WO-US1330.  
 PR 20-NOV-1998; 98WO-US24855.  
 PR 25-NOV-1998; 98WO-US25120.  
 PR 01-DEC-1998; 98WO-US325105.  
 PR 08-MAR-1999; 99WO-US05028.  
 PR 02-JUN-1999; 99WO-US12252.  
 PR 01-SEP-1999; 99WO-US0111.  
 PR 08-SEP-1999; 99WO-US20594.  
 PR 15-SEP-1999; 99WO-US21090.  
 PR 15-SEP-1999; 99WO-US2157.  
 PR 30-NOV-1999; 99WO-US04342.  
 PR 01-DEC-1999; 99WO-US28313.  
 PR 02-DEC-1999; 99WO-US28301.  
 PR 20-DEC-1999; 99WO-US31999.  
 PR 05-FAN-1999; 2000WO-US00219.  
 PR 18-FEB-2000; 2000WO-US04411.  
 PR 18-FEB-2000; 2000WO-US04412.  
 PR 01-MAR-2000; 2000WO-US04414.  
 PR 02-MAR-2000; 2000WO-US05601.  
 PR 09-MAR-2000; 2000WO-US0841.  
 PR 20-MAR-2000; 2000WO-US07377.  
 PR 30-MAR-2000; 2000WO-US04339.  
 PR 15-MAY-2000; 2000WO-US043358.  
 PR 17-MAY-2000; 2000WO-US13705.  
 PR 22-MAY-2000; 2000WO-US14042.  
 PR 02-JUN-2000; 2000WO-US14941.  
 PR 11-AUG-2000; 2000WO-US15264.  
 PR 23-AUG-2000; 2000WO-US23522.

CC Novel secreted and transmembrane polypeptide for modulating biological activity of cell expressing the polypeptide, for identifying agonists or antagonists of polypeptide, and as molecular weight markers -  
 XX DR WPI; 2003-328482/31.  
 XX N-PDBB; ACA60494.  
 XX (GETH ) GENENTECH INC.

XX PI Ashkenazi AJ, Baker KP, Botstein DA, Desnoyers L, Eaton DL,  
 PI Ferrara N, Fong S, Gao W, Gerber H, Goddard A;  
 PI Godowski PJ, Gurkin AL, Kljavin IJ, Mather JP, Napier MA, Pan J;  
 PI Paoni NF, Roy MA, Stewart TA, Tummas D, Watanabe CK, Williams PN;  
 PI Wood WI, Zhang Z;

XX PT Novel secreted and transmembrane polypeptide for modulating biological activity of cell expressing the polypeptide, for identifying agonists or antagonists of polypeptide, and as molecular weight markers -  
 XX DR WPI; 2003-328482/31.  
 XX N-PDBB; ACA60494.

CC The invention describes an isolated, secreted and transmembrane polypeptide (PP), termed PRO PP or fibroblast growth factor receptor PP (I). (I) is useful for detecting PRO33, PRO37, PRO31, PRO187, PRO337, PRO1411, PRO1096, PRO046, PRO607, PRO603, Fibroblast growth factor receptor (FGFR)-3, FGFR-4, FGFR-2, PRO004, PRO356, PRO650, PRO55 or PRO951 polypeptide, and for linking a bioactive molecule to a cell expressing the above polypeptides. The bioactive molecule, a toxin, radiolabel or an antibody, causes cell death. PRO is useful in assays to identify other proteins or molecules involved in binding interaction. The Polynucleotide (II) encoding (I) is useful in chromosome and gene mapping, in generation of antisense RNA and DNA, for generating transgenic animals or knockout animals which in turn are useful in the development and screening of therapeutically useful reagents, to construct hybridisation probes for mapping the gene which encodes the

Db	181	DGHPVITPSRASESASSDGERPVITPSRASESASSDGP	240	Qy	266	PGASDIDLIPTEGVKAASSTDPPALPSTEAKPHITEVTASAETLSTAGTTESSAAPHATV	325
Qy	222	HVITPWSPGDVTLIAELAVTVNTLEVINSITELETTSSIPASDDLDIPIPEGVA	281	Db	313	PGASDIDLIPTEGVKAASSTDPPALPSTEAKPHITEVTASAETLSTAGTTESSAAPHATV	372
Db	241	HVITPWSPGDVTLIAELAVTVNTLEVINSITELETTSSIPASDDLDIPIPEGVA	300	Qy	326	GTPLPNTSATERTVTAPEATTSGALTYVSRNPLEEFSALSVETPSVTKVSGAAPVSEA	385
Qy	282	SSTSDDPALPDSTEAKPHITEVTASAETLSTAGTTESSAAPHATVGPPLPNTSATREVA	341	Db	373	GTPLPNTSATERTVTAPEATTSGALTYVSRNPLEEFSALSVETPSVTKVSGAAPVSEA	432
Db	301	SSTSDDPALPDSTEAKPHITEVTASAETLSTAGTTESSAAPHATVGPPLPNTSATREVA	360	Qy	386	GSAVGKTMISFGASSSNSPSEALKNTFTPSETPTMDIATGPFPFTRDPLPSVPPTTN	445
Qy	342	PGATTLSGALTVYVSRNPLETSALSVETPSVTKVSGAAPVSIERGSAVGKTMISFGASSA	401	Db	433	GSAVGKTMISFGASSSNSPSEALKNTFTPSETPTMDIATGPFPFTRDPLPSVPPTTN	492
Db	361	PGATTLSGALTVYVSRNPLETSALSVETPSVTKVSGAAPVSIERGSAVGKTMISFGASSA	418	Qy	446	SSRGNTNSTLAKITTSAKITMCKP	467
Qy	402	SYSPSEALKNTFPTPSERTPTMDIATKGPRPTSDPLPSVPTTNSRRGTNSTLAKITTS	461	Db	493	SSRGNTNSTLAKITTSAKITMCKP	514
Db	419	SYSPSEALKNTFPTPSERTPTMDIATKGPRPTSDPLPSVPTTNSRRGTNSTLAKITTS	478				
Qy	462	KTTMKPQOPRPRLPGRPQT	482				
Db	479	KTTMKPQPTATP-TTATRPRPT	498				
Qy	492P218	PRELIMINARY;	517 AA.				
Db		PRELIMINARY;	517 AA.				
AC	Q9P218	Q9P218; 15, Created)					
DT	01-OCT-2000	(TREMBrel. 15, Last sequence update)					
DT	01-OCT-2000	(TREMBrel. 15, Last annotation update)					
DE		Hypothetical protein KIAA1359 (Fragment)					
GN	KIAA1359						
OS	Homo sapiens (Human)						
OC	Homo sapiens; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.						
OC	Homo sapiens; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.						
OX	NCBI_TaxID=3606;						
RN	[1]						
RP	SEQUENCE FROM N.A.						
RC	TISSUE=Embryo;						
RC	Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y., Nishikawa T., Nagai K., Sugano S., Shiratori A., Sudo H., Kondo H., Sugawara M., Wagsumma M., Hosoi T., Kaku Y., Kodaira H., Ono Y., Takiguchi S., Takahashi M., Chiba Y., Ishida S., Murakawa K., Ono Y., Saito K., Watanabe S., Kimura K., Murakami K., Ishii S., Kawai Y., Yamamoto J., Wakamatsu A., Nakamura Y., Nagahara K., Masuho Y., Ninomiya K., Iwayanagi T.; RT						
RC	RT	"NEDO human cDNA sequencing project"; RI					
RC	RI	Submitted (MSB-2001) to the EMBL/GenBank/DBJ databases; DR					
RC	EMBL_AK027314; BAB5505.1; -						
RC	KW	Hypothetical protein.					
Medline	20011126; PubMed=16718198;						
RA	Nagase T., Kikuno R., Ishikawa K., Hiroseawa M., Ohara O.; RT						
RA	RT	The complete coding sequences of 150 new cDNA clones from brain which code for large proteins in vitro.";					
RA	RA	DNA Res 7:55-73 (2000); EMBL_AB037780; BAA92597.1; -					
RA	RT	Hypothetical protein.					
RA	RT	HYPOTER_1					
RA	RT	SEQUENCE 517 AA; 52332 MW;	6D14ABA896221DFF CRC64;	Qy	1	MGC1WGLALPFFEFCWEGVGSAGSPSTRRADTMTDDTEVPMILAPGHAALEQTQL	60
RA	RT	Score 2146.5; DB 4; Length 517;		Db	1	MGC1WGLALPFFEFCWEGVGSAGSPSTRRADTMTDDTEVPMILAPGHAALEQTQL	60
RA	RT	Best Local Similarity 87.5%; Pred. No. 1.e-10;		Qy	61	SAETSSRASPAGPPIEAATGKRISSARETSFTSPNFMVLATSVETSAASGPBP	120
RA	RT	Matches 439; Conservative 2; Mismatches 4;		Db	61	SAETSSRASPAGPPIEAATGKRISSARETSFTSPNFMVLATSVETSAASGPBP	120
RA	RT	Indels 57; Gaps 1;		Qy	121	GAGMTVQTITGSDEEAFTLTDSSRBEAATLTMDILTAHTSTAKGLSESSASS	180
RA	RT	23 SSAGPSTRRADTMTDDTEVPMILATSAETSSRASPAGPIPEAETrG 82		Db	121	GAGMTVQTITGSDEEAFTLTDSSRBEAATLTMDILTAHTSTAKGLSESSASS	180
RA	RT	13 TNGPSTARADTMTDDTEVPMILATSAETSSRASPAGPIPEAETrG 72		Qy	181	DGPHPVITPRASESASSDGPHPHVTTPRASESASSDGPHPHVTTPS	240
RA	RT	83 AKRISPARTRTSKTSRPMVLATSVTSAASGPAGHTVQITGSPEEALFD 142		Db	181	DGPHPVITPRASESASSDGPHPHVTTPRASESASSDGPHPHVTTPS	240
RA	RT	73 ARRISPARTRTSKTSRPMVLATSVTSAASGPAGHTVQITGSPEEALFD 132		Qy	241	ALVTVINIEVINCSTIEETTSSPGASDGPHPHVTTPRASESASSDGPHPHVTTPS	300
RA	RT	143 LCDDSSEPAKLTMDILTAHTSTEAKLKS-----173		Db	241	ALVTVINIEVINCSTIEETTSSPGASDGPHPHVTTPRASESASSDGPHPHVTTPS	300
RA	RT	133 LCDDSSEPAKLTMDILTAHTSTEAKLKS-----192		Qy	301	TEVTASAEITLSTAGTTESSAAPHATVGPLPTNSATEREVATGATTGALTVSRNPLE	360
RA	RT	174 -----SESSASSDGPHPVITPRASESASSDGPHPHVTTPRASESASSDGPHPHVTTPS	205	Db	301	TEVTASAEITLSTAGTTESSAAPHATVGPLPTNSATEREVATGATTGALTVSRNPLE	311
RA	RT	193 HPVITPSRASESASSDGPHPVITPRASESASSDGPHPHVTTPRASESASSDGPHPHVTTPS	252				
RA	RT	206 ITPSRASESASSDGPHPVITPRASESASSDGPHPHVTTPRASESASSDGPHPHVTTPS	265				
RA	RT	253 ITPSRASESASSDGPHPVITPRASESASSDGPHPHVTTPS	312				

Tue Jan